STATE AND CONSUMER SERVICES AGENCY DEPARTMENT OF CONSUMER AFFAIRS ARNOLD SCHWARZENEGGER, GOVERNOR

# WHOLESALER DANGEROUS DRUGS & DANGEROUS DEVICES SELF- ASSESSMENT

All legal references used throughout this self-assessment form are explained on page 18.

All references to "drugs" throughout this self-assessment refer to dangerous drugs and dangerous devices as defined in Business & Professions Code (B & P) section 4022. (http://www.pharmacy.ca.gov/laws\_regs/lawbook.pdf).

Wholesaler Name				
Address				
Phone				
Wholesaler E-mail address (op	tional)			
Ownership: Please mark one				
sole owner	partnership	<b>c</b> corporation	LLC LLC	
non- licensed owne	r Cther	(please specify)		
CA Wholesaler Permit #		Expiration Date		
Other Permit #		Expiration Date_		
DEA Registration #		Expiration Date_		
Date of most recent DEA Inven	ntory			
Hours: Daily	Sat	Sun		_ 24 Hours
Designated representative-in-c	harge (DRIC) /	pharmacist (RPH)		
DRIC License # / RPH License # Expiration Date				

### Licensed Wholesaler Staff (designated representative (DR), pharmacist):

1	DR#/RPH#	Exp. Date	
2	DR#/RPH#	Exp. Date	
3	DR#/RPH#	Exp. Date	
4	DR#/RPH#	Exp. Date	
5	DR#/RPH#	Exp. Date	
6	DR#/RPH#	Exp. Date	
7	DR#/RPH#	Exp. Date	
8	DR#/RPH#	Exp. Date	
9	DR#/RPH#	Exp. Date	
10	DR#/RPH#	Exp. Date	

Please mark the appropriate box for each question. If "NO," enter an explanation on the "CORRECTIVE ACTION OR ACTION PLAN" lines at the end of the section. If more space is needed, add additional sheets.

1. Ownership	p/Location
Yes No N/A	Review the current wholesaler permit for this business. Are the listed owners correct and is the listed address correct? If not, please indicate discrepancy. If either is incorrect, notify the board in writing immediately. (B & P 4160[a][c][f]) Attach a copy of the notification letter to the board to this document.
	Have you established and do you maintain a list of officers, directors, managers and other persons in charge of drug distribution, handling and storage? The list must contain a summary of the duties and qualifications for each job listed. (CCR 1780[f][3]) <b>Please attach a copy of the list to this document.</b> (This list should be dated.)
-	request, the owner must provide the board with the names of the owners, managers s and a brief statement of the capacity in which they are employed. (B & P 4082)
CORRECTIV	TE ACTION OR ACTION PLAN
2. Facility	Premises, fixtures and equipment:
Yes No N/A	Are clean and orderly Are well ventilated Are free from rodents and insects Are adequately lit Have plumbing in good repair Have temperature & humidity monitoring to assure compliance with USP Standards. (The standards for various drugs may differ, see USP 1990 22 <sup>nd</sup> Edition) (CCR 1780[b])
	Is there a quarantine area for outdated, damaged, deteriorated, or misbranded drugs, drugs with the outer or secondary seal broken, partially used containers, or any drug returned under conditions that cast doubt on the drugs safety, identity, strength, quality or purity? (CCR 1780[e])

Yes No N/A	Are dangerous drugs and dangerous devices stored in a secured and locked area? (CCR 1780[a])
	Is access to areas where dangerous drugs are stored limited to authorized personnel? (CCR 1780[c])
List personne	l with keys to the area(s) where drugs are stored (list by name or job title):
Yes No N/A	Does this business operate only when a designated representative or pharmacist is on the premises? (CCR 1781)
	The wholesale premises is equipped with the following specific security features: There is an alarm to detect after-hours entry. (CCR 1780[c][1]). The outside perimeter of the building is well lit (CCR 1780[c][3]). The security system provides protection against theft and diversion including tampering with computers and or electronic records. (CCR 1780[c][2]).
Explain how y	your security system complies with these requirements.
Yes No N/A	Is this business a "reverse distributor", that is, does the business act as an agent for pharmacies, drug wholesalers, manufacturers and others, by receiving, inventorying and managing the disposition of outdated or nonsalable drugs? (B & P 4040.5)
CORRECTIV	TE ACTION OR ACTION PLAN
	are specific requirements for wholesaling controlled substances – these additional are in Section 11 of this document.

## 3. Designated Representative-in-Charge / Owner Responsibilities Yes No N/A The owner and the designated representative-in-charge are both equally responsible for maintenance of the records and inventory. (B & P 4081[b]) Is the designated representative-in-charge responsible for the wholesaler's compliance with all state and federal laws for the wholesale distribution of drugs? The designated representative-in-charge may be a pharmacist. (B & P 4160[d]) The owner must notify the board within 30 days of termination of the designated representative-in-charge or pharmacist. (B & P 4305.5[a]) The owner must identify and notify the board of the appointment of a new designated representative-in-charge within 30 days of the termination of the former designated representative-in-charge. (B & P 4160[d], 4331[c]) The appropriate form for this notification is a "Change of Designated Representativein-Charge," which is available on the board's website. The designated representative-in-charge who ends his or her employment at a wholesaler, must notify the board within 30 days. (B & P 4305.5[c], 4101[b]). This notification is in addition to that required of the owner. CORRECTIVE ACTION OR ACTION PLAN \_\_\_\_\_ 4. Designated Representative/Pharmacist Yes No N/A If a designated representative or pharmacist changes his/her name or personal address of record, he/she must notify the board in writing within 30 days. (B & P 4100, 1704) CORRECTIVE ACTION OR ACTION PLAN 5. Ordering Drugs by this Business for Future Sale/Transfer or Trade Yes No N/A Are drugs ordered only from a business licensed by this board or from a licensed manufacturer? (B & P 4163[b], 4169)

Yes No N/A	If drugs are returned to your premises by a business that originally purchased the drugs from you, do you document the return with an acquisition record for your business and a disposition record for the business returning the drugs? (B & P 4081, 4332)	
CORRECTIVE ACTION OR ACTION PLAN		
	are in Section 11 of this document.	
6. Receipt of	Drugs by this Business	
Yes No N/A	When drugs are received by your business, are they delivered to the licensed wholesale premises, and received by and signed for only by a designated representative or a pharmacist? (B & P 4059.5[a])	
	When drugs are received by your business, are the outside containers visibly inspected to identify the drugs and prevent acceptance of contaminated drugs by detecting container damage? (CCR 1780[d][1])	
CORRECTIV	E ACTION OR ACTION PLAN	
	are in Section 11 of this document.	
7. Drug Stock	<b>C</b>	
Yes No N/A	Is all drug stock open for inspection during regular business hours? (B & P 4081[a])	
	Are all drugs you order maintained in a secure manner at your licensed wholesale premises? You cannot order, obtain or purchase drugs that you are not able to store on your licensed premises. (B & P 4167)	
	Do all drugs you sell conform to the standards and tests for quality and strength provided in the latest edition of United States Pharmacopoeia or Sherman Food Drug and Cosmetic Act? (B & P 4342[a])	

Yes No N/A	Do all drug containers you store on your premises have a manufacturer's expiration date? Any drug without an expiration date is considered expired and may not be distributed. (CCR 1718.1)
	Are outdated, damaged, deteriorated or misbranded drugs held in a quarantine area physically separated from other drugs until returned to the supplier or sent for destruction? (CCR 1780[e], CFR 1307.21)
	Are drugs with the outer or secondary seal broken, or partially used or returned drugs held in a quarantine area and physically separated from other drugs until returned to the supplier or sent for destruction? (CCR 1780[e], CFR1307.21)
	When the conditions under which drugs were returned to your premises cast doubt on the drugs' safety, identity, strength, quality or purity, are the drugs quarantined and either returned to your supplier or destroyed? If testing or investigation proves the drugs meet USP standards, the drugs may be returned to normal stock. (CCR 1780[e], CFR 1307.21)
CORRECTIV	VE ACTION OR ACTION PLAN
requirements	are specific requirements for wholesaling controlled substances – these additional are in Section 11 of this document.  ansfer of Drugs by this Business
Yes No N/A	Are drugs sold only to businesses or persons licensed by this board, licensed by a prescriber board, licensed as a manufacturer, or to a licensed health care entity authorized to receive drugs?
Describe how B & P 4169)	you verify a business or person is appropriately licensed. (B & P 4059.5[a] [b][d],
List any busir list above:	nesses or individuals that order drugs from you that are not licensed according to the

Yes No N/A	Are drugs only furnished by your business to an authorized person? (B & P 4163[a]) Note: An authorized person can be a business or natural person.
	Does your business only receive drugs from a pharmacy if: the pharmacy originally purchased the drugs from you? your business is a "reverse distributor"? the drugs are needed to alleviate a shortage? (and only a quantity sufficient to alleviate a specific shortage). (B & P 4126.5[a])
	Are all drugs that are purchased from another business or that are sold, traded or transferred by your business:
	transacted with a business licensed with this board as a wholesaler or
	pharmacy? free of adulteration as defined by the CA Health & Safety Code section 111250?
	free of misbranding as defined by CA Health & Safety Code section 111335?
	confirmed to not be beyond their use date (expired drugs)? (B & P 4169)
	dents where adulterated, misbranded or expired drugs were purchased, sold, traded d by this business in the past 2 years.
	If your business sells, transfers, or delivers dangerous drugs or devices outside of California, either to another state within the United States or a foreign country, do
Yes No N/A	you: comply with all CA pharmacy laws related to the distribution of drugs? comply with the pharmacy law of the receiving state within the United States?
	comply with the statues and regulations of the Federal Food and Drug Administration and the Drug Enforcement Administration relating to the wholesale distribution of drugs?
	comply with all laws of the receiving foreign country related to the wholesale distribution of drugs?
	comply with all applicable federal regulations regarding the exportation of dangerous drugs?
	w you determine a business in a foreign country is authorized to receive dangerous gerous devices. (B & P 4059.5[e])
-	

Yes No N/A	When you are not an authorized distributor for a drug, a pedigree must accompany the product when sold, traded, or transferred (Prescription Drug Marketing Act of 1987). Effective January 1, 2009, an electronic pedigree must accompany all drugs (B & P 4163), even those for which your business is an authorized distributor.
	If preferentially priced drugs are sold by your business, that sale complies with the Prescription Drug Marketing Act of 1987 and CA Pharmacy Law. (B & P 4380)
	Does your business' advertisements for dangerous drugs or devices contain false, fraudulent, misleading or deceptive claims? (B & P 4341, B & P 651, CCR 1766)
	Do you offer or receive any rebates, refunds, commissions or preferences, discounts or other considerations for referring patients or customers? If your business has any of these arrangements, please list with whom. (B & P 650)
Yes No N/A	Does your business sell dangerous drugs or devices to the master or first officer of an ocean vessel, after your business has received a written prescription? If so, describe how you comply with the ordering, delivery and record keeping requirements for drugs including controlled substances, and the requirement to notify the board of these sales. (B & P 4066, CFR 1301.25)
CORRECTIV	E ACTION OR ACTION PLAN
	are specific requirements for wholesaling controlled substances – these additional are in Section 11 of this document.
9. Outgoing S	Shipments of Drugs
Yes No N/A	Before you ship drugs to a purchaser, do you inspect the shipment to assure the drugs were not damaged while stored by your business? (CCR 1780[d][2])

Yes No N/A	Does your business use a common carrier (a shipping or delivery company — UPS, US Mail, FedEx, DHL) for delivery of drug orders to your customers? (B & P 4166[a])	
List the common carriers (shipping or delivery companies) you use.		
CORRECTIV	E ACTION OR ACTION PLAN	
	are in Section 11 of this document.	
10. Delivery	of Drugs	
Yes No N/A	Are all drugs ordered by a pharmacy or another wholesaler delivered to the address of the buyer's licensed premises and signed for and received by a pharmacist or designated representative where allowed? (B & P 4059.5[a])	
	Are all drugs ordered by a manufacturer or prescriber delivered to the manufacturer's or prescriber's licensed business address and signed for by a person duly authorized by the manufacturer or prescriber? (B & P 4059[d])	
	All drugs delivered to a hospital are delivered either to the pharmacy premises or to a central receiving area within the hospital. (B & P 4059.5[c])	
	If drugs are delivered to a pharmacy when the pharmacy is closed and a pharmacist is not on duty, documents are left with the delivery in the secure storage facility, indicating the name and amount of each dangerous drug delivered. (B & P 4059.5[f])	
CORRECTIV	E ACTION OR ACTION PLAN	
11. Controlle	d Substances	
Yes No N/A	Are there effective controls to prevent theft or diversion of controlled substances? (CFR 1301.71)	

Yes No N/A	Are DEA requirements for storage of Schedule II controlled substances being met? (specific requirements are listed in CFR 1301.72[a])
	Are DEA requirements for storage of Schedule III controlled substances being met? (specific requirements are listed in CFR 1301.72[b])
	Is a DEA inventory completed by your business every two years for all schedules (II - V) of controlled substances? (CFR 1304.11[a][c][e])
	Is the biennial record of the DEA inventory required for Schedule II – V controlled substances conducted every 2 years, retained for 3 years? (CFR 1304.11, CCR 1718, 1780(f)[2])
	Has the person within your business who signed the original DEA registration, or the last DEA registration renewal, has created a power of attorney for each person allowed to order Schedule II controlled substances for this business? (CFR 1305.05)
List the indiv substances.	iduals at this location authorized by power of attorney to order controlled
Yes No N/A	Does your business follow employee-screening procedures required by DEA to assure the security of controlled substances? (CFR 1301.90)
	If any employee of this business possesses, sells, uses or diverts controlled substances, in addition to the criminal liability you must evaluate the circumstances of the illegal activity and determine what action you should take against the employee. (CFR 1301.92)
	Are all controlled substances purchased, sold or transferred by your business, done so for legitimate medical purposes? (H & S 11153.5[a][b][c])
	If your business distributes controlled substances through an agent (i.e. detail person), do you have adequate security measures in place to prevent theft or diversion of those controlled substances (CFR 1301.74[f])
	If a person attempts to purchase controlled substances from your business and the person is unknown to you, you make a good faith effort to determine the person (individual or business) is appropriately licensed to purchase controlled substances. (CFR 1301.74 [a])

_	your business determines an unknown business or individual is appropriately urchase controlled substances
Yes No N/A	If your business uses a common carrier to deliver controlled substances, your business determines the common carrier has adequate security to prevent the theft or diversion of controlled substances.(CFR 1301.74[f])
	If your business uses a common carrier to deliver controlled substances, are the shipping containers free of any outward indication that there are controlled substances within, to guard against storage or in-transit theft? (CFR 1301.74[e])
	Are all Schedule II controlled substances ordered from your business using a fully completed DEA 222 order form? (CFR 1305.03, 1305.06)
	When your business fills orders for Schedule II controlled substances, is the date filled and the number of containers filled recorded on copies 1 and 2 of DEA 222 from? Is copy 1 retained and copy 2 sent to DEA at the close of the month the controlled substance order was filled? (CFR 1305.13 [b])
	If a Schedule II controlled substance order cannot be filled, does your business return copy 1 and 2 of the DEA 222 order form to the buyer with a letter indicating why the order could not be filled? (CFR 1305.15)
	When your business partially fills Schedule II controlled substances, is the balance provided within 60 days of the date of the order form? After the final partial filling, is copy 1 retained in your files and copy 2 of the completed DEA 222 order form sent to DEA by the close of that month? (CFR 1309.13[b])
	For all Schedule II controlled substances received by your business, is copy 3 of the DEA 222 order form completed by writing in for each item received, the date received and the number of containers received? (CFR 1305.13[e])
	Does your business use the online CSOS secure transmission system offered by the Drug Enforcement Administration in place of a paper DEA 222 Form for Schedule II controlled substances?
	Does your business follow the procedure outlined by DEA to obtain Schedule II controlled substances when the original DEA 222 order form is lost or stolen? (CFR 1305.16(a))
	Are all records of purchase and sale for all schedules of controlled substances for your business kept on your licensed business premises for 3 years from the

X/ N/ N/A	making? (B & P 4081, CCR 1718, CFR 1305.09[d], 1305.17[a] [b], and H & S 11252, 11253, 1304.03)
Yes No N/A	Are records of Schedule II controlled substances stored separate from all others? (CFR 1304.04 [f][1])
	Are records for Schedule III-V controlled substances stored so that they are easily retrievable? (CFR 1304.04 [f][2])
	Before your business distributes carfentanil etorphine HCL and or diprenorphine, do you contact the DEA to determine the person (individual or business) is authorized to receive these drugs? (CFR 1301.75[g], 1305.16[b])
	Do you separate records for the sale of carfentanil etorphine hydrochloride and or diprenorphine from all other records? (CFR 1305.16)
	Does the owner of your business notify the DEA, on a DEA 106 form, of any theft or significant loss of controlled substances upon discovery of the theft? (CFR 1301.74[c])
	Does the owner of your business notify the board of any loss of controlled substances within 30 days of discovering the loss? (CCR 1715.6)
	and Procedures
	Does this business maintain and adhere to policies and procedures for:
Yes No N/A	Receipt of drugs? Security of drugs? Storage of drugs? (including maintaining records to document proper
	storage) Inventory of drugs? (including correcting inaccuracies in inventories) Distributing drugs? Identifying, recording and reporting theft or losses? Correcting errors?
	Physically quarantining and separating: returned, damaged, outdated, deteriorated, misbranded or adulterated drugs?
	drugs that have been partially used? drugs where the outer or secondary seals on the container have been broken?
	drugs returned to your business, when there is doubt about the safety, identity, strength, quality, or purity of the drug?

Yes No N/A	drugs where the conditions of return cast doubt on safety, identity, strength, quality or purity? (CCR 1780[e][f])
CORRECTIV	VE ACTION OR ACTION PLAN
13. Training	
Yes No N/A	Is training and experience provided to all employees to assure all personnel comply with all licensing requirements? (CCR 1780[f][4])
List the types training.	s of training you have provided to staff in the last calendar year and the dates of that
CORRECTIV	VE ACTION OR ACTION PLAN
14. Dialysis	Drugs
Yes No N/A	Does your business provide dialysis drugs directly to patients, pursuant to a prescription? (B & P 4054) (4059[c]) If so, please complete the next 4 questions, if not proceed to Section 15.
	Do home dialysis patients complete a training program provided by a dialysis center licensed by Department of Health Services? Prescriber must provide proof of completion of this training to your business. (B & P 4059[d])
	Do you have written or oral orders for authorized dialysis drugs for each dialysis patient being serviced. Are such orders received by either a designated representative or a pharmacist? Note: refill orders cannot be authorized for more than 6 months from the date of the original order. (CCR 1787[a][b][c])
	Does your business provide an "expanded invoice" for dialysis drugs dispensed directly to the patient including name of drug, manufacturer, quantities, lot number, date of shipment, and name of the designated representative or pharmacist responsible for distribution? A copy of the invoice must be sent to the prescriber, the patient and a copy retained by this business. Upon receipt of drugs,

Yes No N/A	the patient or patient agent must sign for the receipt for the drugs with any irregularities noted on the receipt. (CCR 1790)			
	Is each case or full shelf package of the dialysis drugs dispensed labeled with the patient name and the shipment? Note that additional information as required is provided with each shipment. (CCR 1791)			
CORRECTIVE ACTION OR ACTION PLAN				
15 December 1	Cooning Dequinoments			
15. Record N	Ceeping Requirements			
Yes No N/A	Does your business' sales record for drugs include date of sale, your business name and address, the business name and address of the buyer, and the names and quantities of the drugs sold? (B & P 4059[b])			
	Are purchase and sales records for all transactions retained on your licensed premises for 3 years from the date of making? (B & P 4081[a], 4105[c], 4081, 4332, 4059.5[a]) Note: A drug pedigree is considered to be a part of the records of purchase and sale and must be retained for three years from the making.			
	Are all purchase and sales records retained in a readily retrievable form? (B & P 4105[a])			
	Is a current accurate inventory maintained for all dangerous drugs? (B & P 4081, 4332, 1718)			
	If you temporarily remove purchase or sales records from your business, does your business retain on your licensed premises at all times, a photocopy of each record temporarily removed? (B & P 4105[b])			
	Are required records stored off-site only if a board issued written waiver has been granted?			
-	ss has a written waiver, write the date the waiver was approved and the off-site the records are stored below. (CCR 1707[a])			
Date	Address			
Yes No N/A	Is an off-site written waiver in place and is the storage area secure from unauthorized access? (CCR 1707[b][1])			
	If an off-site written waiver is in place, are the records stored off-site retrievable within 2 business days? (CCR 1707[b][2])			

Yes No N/A	Can the records that are retained electronically be produced immediately in hard copy form by any designated representative, if the designated representative-incharge is not present? (B & P 4105[d])
	Are records of training provided to employees to assure compliance with licensing requirements, retained for 3 years? (CCR 1780[f][4])
	Has this licensed premises, or the designated representative-in-charge or pharmacist, been cited, fined or disciplined by this board or any other state or federal agency within the last 3 years? If so list each incident with a brief explanation (B & P 4162[a][4]):
Yes No N/A	Has the licensed premises received any orders of correction from this board? A copy of the order and the corrective action plan must be on the licensed premises for 3 years. (B & P 4083)
	Has this business received a letter of admonishment from this board? A copy must be retained on the premises for 3 years from the date of issue. (B & P 4315[e])
	If this business dispenses dialysis drugs directly to patients, are the prescription records retained for 3 years, including refill authorizations and expanded invoices for dialysis patients? (CCR 1787[c], 1790)
CORRECTIV	E ACTION OR ACTION PLAN
	are specific requirements for wholesaling controlled substances – these additional are in Section 11 of this document.
16. Reporting	g Requirements to the Board
Yes No N/A	A designated representative-in-charge who terminates employment at this business, must notify the board within 30 days of the termination (B & P 4101[b], 4305.5[c].
	The owner must report to the board within 30 days the termination of the designated representative-in-charge or pharmacist (B & P 4305.5[a])

Yes No N/A	The owner must report to the board within 30 days of discovery, any loss of controlled substances, including amounts and strengths of the missing drugs. (CCR 1715.6)
	The owner must notify the DEA, on a DEA form 106, any theft or significant loss of controlled substances upon discovery. (CFR 1301.74[c])
	Do your employees know about their obligation to report any known diversion or loss of controlled substances to a responsible person within your business? (CFR 1301.91)
	The owner must notify the board within 30 days of any change in the beneficial ownership of this business. (B & P 4201[i], CCR 1709[b])
	When called upon by the board, your business can report all sales of dangerous drugs or controlled substances subject to abuse. (B & P 4164[a])
	Effective January 1, 2006 your business will develop and maintain a tracking system for individual sales of dangerous drugs at preferential or contract prices to pharmacies that primarily or solely dispense prescription drugs to patients of long-term care facilities. Your system must:  1. identify pharmacies that primarily or solely dispense prescription drugs to patients of long term care facilities  2. identify purchases of any dangerous drugs at preferential or contract prices  3. identify current purchases that exceed prior purchases by 20 percent over the previous 12 calendar months. (B & P 4164[b])
	I understand that this wholesaler license is not transferable to a new owner. A change of ownership must be reported to this board, as soon as the parties have agreed to the sale. Before the ownership actually changes, an additional application for a temporary permit must be submitted to the board if the new owner wants to conduct business while the board is processing the change of ownership application and until the new permanent permit is issued. A company cannot transfer the ownership of the business via a contract with another individual or business, without the board's approval (B & P 4201[g])
	The owner of this business must immediately notify the board in writing if any assignment is made for the benefit of creditors, if the business enters into any credit compromise arrangement, files a petition in bankruptcy, has a receiver appointed, or enters into liquidation or any other arrangement that might result in the sale or transfer of drugs. (CCR 1705)
	If this business is discontinued, the owner must notify the board in writing before the actual discontinuation of business. (CCR 1708.2). If the business holds a DEA registration, the owner must notify the DEA promptly of the discontinuation of business and all unused DEA 222 order forms must be returned to the DEA. (CFR 1301.52[a], 1305.14)

CORRECTIVE ACTION OR ACTION PLAN
17. Additional Licenses/Permits Required
List all licenses and permits required to conduct this business, including local business licenses, wholesale licenses held in other states, permits or licenses required by foreign countries or other entities (B & P 4107, CFR 1305.11[a], B & P 4059.5[e])
DESIGNATED REPRESENTATIVE-IN-CHARGE / PHARMACIST CERTIFICATION:
I, (please print), DRIC# / RPH # hereby certify that I have completed the self-assessment of this wholesale business of which I am the designated representative-in-charge (DRIC) / pharmacist (RPH). I understand that all responses are subject to verification by the Board of Pharmacy. I further state under penalty of perjury that the information contained in this self-assessment form is true and correct.
Signature Date Date

#### **Legal References**

All references to California Business & Professions Code (B & P) are Chapter 9, Division 2 unless otherwise specified (<a href="http://www.pharmacy.ca.gov/laws\_regs/lawbook.pdf">http://www.pharmacy.ca.gov/laws\_regs/lawbook.pdf</a>).

All references to California Code of Regulations (CCR) are to Title 16 unless otherwise specified (<a href="http://www.pharmacy.ca.gov/laws\_regs/lawbook.pdf">http://www.pharmacy.ca.gov/laws\_regs/lawbook.pdf</a>).

All references to California Health & Safety Code (H & S) are to Division 10, Uniform Controlled Substances Act (<a href="http://www.pharmacy.ca.gov/laws\_regs/lawbook.pdf">http://www.pharmacy.ca.gov/laws\_regs/lawbook.pdf</a>) or Division 104, Part 5, Sherman Food, Drug and Cosmetic Laws <a href="http://www.dhs.ca.gov/fdb/PDF/Sherman%202006.PDF">http://www.dhs.ca.gov/fdb/PDF/Sherman%202006.PDF</a>

All references to United States Code of Federal Regulations (CFR) are Title 21, Chapter II Part 1300, Drug Enforcement Administration, Food and Drugs and codified Controlled Substances Act (CSA) (<a href="http://www.deadiversion.usdoj.gov/21cfr/index.html">http://www.deadiversion.usdoj.gov/21cfr/index.html</a>).

#### California Board of Pharmacy

1625 N. Market Blvd., Suite N219 Sacramento CA 95834 (916) 574-7900 fax: (916) 574-8618 www.pharmacy.ca.gov

#### California Pharmacy Law may be obtained

by contacting: Law Tech 1060 Calle Cordillera, Suite 105 San Clements CA 92673 (800) 498-0911 Ext. 5 www.lawtech-pub.com

#### **Pharmacist Recovery Program**

(800) 522-9198 (24 hours a day)

#### **Prescriber Boards:**

#### **Medical Board of California**

1426 Howe Avenue, Suite 54 Sacramento CA 95825 (800) 633-2322 (916) 263-2499 fax: (916) 263-2387 http://www.mbc.ca.gov

#### **Dental Board of California**

1432 Howe Ave. #85 Sacramento, CA 95825 (916) 263-2300 fax: (916) 263-2140 http://www.dbc.ca.gov

#### **Board of Registered Nursing**

1625 N. Market Blvd., Suite N217 Sacramento, CA 95834 (916) 322-3350 fax: (916) 574-8637 http://www.rn.ca.gov/

#### **Board of Optometry**

2420 Del Paso Road, Suite 255 Sacramento, CA 95834 (916) 575-7170 fax: (916) 575-7292 http://www.optometry.ca.gov/

#### Osteopathic Medical Board of California

2720 Gateway Oaks Drive, #350 Sacramento, CA 95833 (916) 263-3100 fax: (916) 263-3117 http://www.ombc.ca.gov

#### **Physician Assistant Committee**

1424 Howe Avenue, #35 Sacramento, CA 95825 (916) 561-8780 fax: (916) 263-2671 http://www.physicianassistant.ca.gov

#### **Board of Podiatric Medicine**

1420 Howe Avenue, #8 Sacramento, CA 95825 (800) 633-2322 (916) 263-2647 fax: (916) 263-2651 http://www.bpm.ca.gov

#### **Veterinary Medical Board**

1420 Howe Avenue, #6 Sacramento, CA 95825 (916) 263-2610 fax: (916) 263-2621 http://www.vmb.ca.gov

#### **Federal Agencies:**

## Food and Drug Administration - Industry Compliance

http://www.fda.gov/oc/industry/centerlinks.html#drugs

# The **Drug Enforcement Administration** may be contacted at:

#### **DEA Website:**

http://www.deadiversion.usdoj.gov

#### **Online Registration – New Applicants:**

http://www.deadiversion.usdoj.gov/drugreg/reg\_apps/onlineforms new.htm

#### **Online Registration - Renewal:**

www.deadiversion.usdoj.gov/drugreg/reg\_apps/onlineforms.htm

#### **Registration Changes (Forms):**

http://www.deadiversion.usdoj.gov/drugreg/chan ge requests/index.html

#### **Online DEA 106 Reporting:**

https://www.deadiversion.usdoj.gov/webforms/app106Login.jsp

#### **Controlled Substance Ordering System**

(CSOS): http://www.deaecom.gov/

#### **DEA Registration Support (all of CA):**

(800) 882-9539

#### **DEA - Los Angeles**

255 East Temple Street, 20th Floor Los Angeles CA 90012 (888) 415-9822 or (213) 621-6960 (Registration) (213) 621-6942 or 6952 (Diversion or Investigation)

#### **DEA - San Francisco**

450 Golden Gate Avenue San Francisco CA 94102 Registration: (888) 304-3251 or

(415) 436-7900

Theft Reports or Diversion: (415) 436-7854

#### **DEA - Sacramento**

4328 Watt Avenue Sacramento CA 95821

Registration: (888) 304-3251 or

(415) 436-7900

Diversion or Investigation: (916) 480-7100

or (916) 480-7250

#### **DEA - Riverside**

4470 Olivewood Avenue Riverside, CA 92501-6210 Registration: (888) 415-9822 or

(213) 621-6960

Diversion or Investigation: (909) 328-6000

or (909) 328-6200

#### **DEA - Fresno**

2444 Main Street, Suite 240

Fresno, CA 93721

Registration: (888) 304-3251 or

(415) 436-7900

Diversion or Investigation: (559) 487-5402

#### **DEA – San Diego and Imperial Counties**

4560 Viewridge Avenue San Diego, CA 92123-1637 Registration: (800) 284-1152

Diversion or Investigation: (858) 616-4100

#### DEA - Oakland

1301 Clay Street, Suite 460N

Oakland, CA 94612

Registration: (888) 304-3251 or

(415) 436-7900

Diversion or Investigation: (510) 637-5600

#### DEA – San Jose

One North First Street, Suite 405

San Jose, CA 95113

Registration: (888) 304-3251 or

(415) 436-7900

Diversion or Investigation: (408) 291-7620

or (408) 291-2631

#### **DEA – Redding**

310 Hensted Drive, Suite 310

Redding, CA 96002

Registration: (888) 304-3251 or

(415) 436-7900

Diversion or Investigation: (530) 246-5043